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                 MEDLINE segment
NEWS 13 DEC 17 MEDLINE and LMEDLINE updated with 2008 MeSH vocabulary
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                 STN Viewer enhanced with full-text patent content
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NEWS 20 JAN 28 USGENE now provides USPTO sequence data within 3 days
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NEWS 24 FEB 20 PCI now available as a replacement to DPCI
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                 applications updated
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                 LPCI now available as a replacement to LDPCI
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50 DOCS

100.0% DONE 646 VERIFIED 307 HIT RXNS INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE \*\*COMPLETE\*\*

BATCH \*\*COMPLETE\*\*

PROJECTED VERIFICATIONS: 11396 TO 14444 PROJECTED ANSWERS: 850 TO 1830

L2 50 SEA SSS SAM L1 ( 307 REACTIONS)

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100.0% DONE 13218 VERIFIED 7572 HIT RXNS 1237 DOCS

SEARCH TIME: 00.00.05

L3 1237 SEA SSS FUL L1 ( 7572 REACTIONS)

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=> s 13

L4 1237 L3

=> s 13 and free radical 1237 L3 1368029 FREE 328527 RADICAL 74182 FREE RADICAL

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(FREE (W) RADICAL)
T<sub>1</sub>5
            13 L3 AND FREE RADICAL
=> s 13 and radical
          1237 L3
        328527 RADICAL
            98 L3 AND RADICAL
L6
=> s 13 and initiator
          1237 L3
         74166 INITIATOR
L7
             4 L3 AND INITIATOR
=> s peroxide? or diazo? or dialkylphenylalkane?
        247525 PEROXIDE?
        112472 DIAZO?
             0 DIALKYLPHENYLALKANE?
        359023 PEROXIDE? OR DIAZO? OR DIALKYLPHENYLALKANE?
T.8
\Rightarrow s 18 and 13
          1237 L3
1.9
            28 L8 AND L3
=> s 19 and 16
             4 L9 AND L6
=> s 17 or 110
             8 L7 OR L10
=> s 111 not py > 2003
       5547651 PY > 2003
             6 L11 NOT PY > 2003
L12
=> d 112 ibib abs hitstr 1-
YOU HAVE REQUESTED DATA FROM 6 ANSWERS - CONTINUE? Y/(N):y
L12 ANSWER 1 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER:
                         2001:228100 CAPLUS
DOCUMENT NUMBER:
                         135:92680
TITLE:
                         Iodine atom transfer addition reaction of 1-iodoalkyl
                         phosphonates to alkenes in the presence of
                         \alpha, \alpha'-azoisobutyronitrile (AIBN):
                         mechanistic aspects
                         Balczewski, Piotr; Mikolajczyk, Marian
AUTHOR(S):
CORPORATE SOURCE:
                         Centre of Molecular and Macromolecular Studies, Polish
                         Academy of Sciences, Lodz, 90-363, Pol.
                         New Journal of Chemistry (2001), 25(4), 659-663
SOURCE:
                         CODEN: NJCHE5; ISSN: 1144-0546
                         Royal Society of Chemistry
PUBLISHER:
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
                         CASREACT 135:92680
OTHER SOURCE(S):
     The mechanistic pathway of the title reaction, which constitutes the 1st
     example of a radical I atom transfer addition reaction of
nonfluorine-containing
     phosphonates, will determine whether 2-iodo-2-methylpropionitrile, 8, can serve
     as a competing I donor with the starting di-Et 1-iodoalkyl phosphonates,
     la,b. The title reaction proceeds with AIBN as the sole radical
     initiator, not requiring poisonous Sn reagents as co-initiators,
     and gave di-Et 3-iodoalkylphosphonates 3a-e (the final products of the
     propagation step, isolated in 59-95% yield), tetramethylsuccinodinitrile,
     9, di-Et methylphosphonate, 4, and tetra-Et ethylenebisphosphonate 5 (all
     termination products, 0-10% yields). The radical character of this
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reaction was demonstrated using TEMPO as a radical trap. 8 (the intermediate of the initiation step), synthesized independently from AIBN and I, caused complete inhibition of the reaction when added to the reaction mixture, indicating that it does not behave as an I donor in the transfer stage, but rather as an inhibitor.

REFERENCE COUNT: 25 THERE ARE 25 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L12 ANSWER 2 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1996:656498 CAPLUS

DOCUMENT NUMBER: 125:328880

TITLE: The Synthesis of Phosphonate Ester Containing

Fluorinated Vinyl Ethers

AUTHOR(S): Pedersen, Scot D.; Qiu, Weiming; Qiu, Zai-Ming; Kotov,

Stefan V.; Burton, Donald J.

CORPORATE SOURCE: Department of Chemistry, University of Iowa, Iowa

City, IA, 52242, USA

SOURCE: Journal of Organic Chemistry (1996), 61(23), 8024-8031

CODEN: JOCEAH; ISSN: 0022-3263

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 125:328880

Three novel perfluorovinyl ethers containing phosphonate ester groups, di-Et 1,1,2,2,3,3,5,6,6-nonafluoro-4-oxa-5-hexenylphosphonate, (EtO) 2P(O) (CF2) 3OCF: CF2 (1), di-Et 1,1,2,2,4,5,5-heptafluoro-3-oxa-4pentenylphosphonate, (EtO)2P(O)(CF2)2OCF:CF2 (2), and di-Et 1,1,2,2,4,5,5,7,8,8-decafluoro-4-trifluoromethyl-3,6-dioxa-7octenylphosphonate, CF2:CFOCF2CF(CF3)O(CF2)2P(O)(OEt)2 (3), have been synthesized. Perfluoro vinyl ethers 1 and 2 were synthesized from Me 4-trifluoroethenoxy-2,2,3,3,4,4-hexafluorobutanoate and Me 3-trifluoroethenoxy-2,2,3,3-tetrafluoropropanoate, resp., while perfluorovinyl ether 3 was synthesized either from 5-trifluoroethenoxy-4trifluoromethyl-3-oxa-1,1,2,2,4,5,5-heptafluoropentylsulfonyl fluoride or Me 6-trifluoroethenoxy-5-trifluoromethyl-4-oxa-2,2,3,3,5,6,6heptafluorohexanoate. The carboxylate esters were converted to the corresponding fluoroalkyl iodides via a free-radical iododecarboxylation. The sulfonyl fluoride was converted to its corresponding fluoroalkyl iodide via iododesulfination. The intermediate iodides were useful precursors for the incorporation of the phosphonic ester groups via a photoreaction with tetra-Et pyrophosphite to produce di-Et fluorophosphonites. The di-Et fluorophosphonites were oxidized to the desired phosphonates, 1, 2, and 3, utilizing hydrogen peroxide as the oxidant. Moderate to good overall yields of perfluorovinyl ethers 1-3 have been achieved.

L12 ANSWER 3 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1993:255278 CAPLUS

DOCUMENT NUMBER: 118:255278

TITLE: Stereocontrolled construction of substituted

pyrrolidines based on intramolecular protodesilylation reaction. Enantiospecific synthesis of (-)-kainic acid

and (+)-allokainic acid from L-serine

AUTHOR(S): Hatakeyama, Susumi; Sugawara, Kazutoshi; Takano,

Seiichi

CORPORATE SOURCE: Pharm. Inst., Tohoku Univ., Sendai, 980, Japan

SOURCE: Journal of the Chemical Society, Chemical

Communications (1993), (2), 125-7 CODEN: JCCCAT; ISSN: 0022-4936

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 118:255278

GΙ

AB Novel stereocontrolled enantiospecific syntheses of (-)-kainic acid (I) and (+)-allokainic acid (II) have been achieved starting from L-serine via two modes of C-2 and C-3 side chain-directed intramol. protodesilylations of 4-(trimethylsilylmethyl)ethylidenepyrrolidines. Thus, the BF3.Et2O-mediated intramol. protodesilylation reaction of silylated pyrrolidine derivative III (TMS = trimethylsilyl, TBDPS = tert-butyldiphenylsilyl) followed by methylation with CH2N and HF cleavage gave 3,4-trans-pyrrolidine IV with complete diastereoselectivity. IV was converted to II in 3 steps. On the other hand, the BF3.Et2O-mediated intramol. protodesilylation reaction of silylated pyrrolidine derivative V under diluted conditions proceeded with opposite diastereoselectivity to give 3,4-cis-pyrrolidine VI and its C-4 epimer in a ratio of 5.3:1. VI was converted to I.

L12 ANSWER 4 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1990:7316 CAPLUS

DOCUMENT NUMBER: 112:7316

TITLE: Stereoselective exocyclic double-bond formation via

vinyl-radical cyclization

AUTHOR(S): Munt, Simon P.; Thomas, Eric J.

CORPORATE SOURCE: Dyson Perrins Lab., Oxford, OX1 3QY, UK SOURCE: Journal of the Chemical Society, Chemical

Communications (1989), (8), 480-2 CODEN: JCCCAT; ISSN: 0022-4936

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 112:7316

GΙ

MeO<sub>2</sub>CHC CO<sub>2</sub>Me

AB A stereoselective route to 2,6-cis-disubstituted 4-methoxycarbonylmethylenetetrahydropyrans, e.g., I, has been developed which uses a vinyl radical cyclization to establish the geometry of the exocyclic double bond, e.g., of MeO2CCH:CRCH2CH2OC(:CH2)CO2Me (R = Br, iodo).

L12 ANSWER 5 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1985:166291 CAPLUS

DOCUMENT NUMBER: 102:166291

ORIGINAL REFERENCE NO.: 102:26137a,26140a

TITLE: Exchange of bromine and chlorine by iodine in

haloaliphatic compounds induced by pentacarbonyliron

AUTHOR(S): Amriev, R. A.; Velichko, F. K.; Freidlina, R. Kh.

CORPORATE SOURCE: Inst. Elementoorg. Soedin., Moscow, USSR

SOURCE: Izvestiya Akademii Nauk SSSR, Seriya Khimicheskaya

(1985), (1), 226-8

CODEN: IASKA6; ISSN: 0002-3353

DOCUMENT TYPE: Journal LANGUAGE: Russian

OTHER SOURCE(S): CASREACT 102:166291

AB Fe(CO)5 initiated the exchange reactions of RX (R = pentyl, Bu, allyl; X = Cl, Br) with I2 or CHI3 to give RI.

L12 ANSWER 6 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1985:45503 CAPLUS

DOCUMENT NUMBER: 102:45503

ORIGINAL REFERENCE NO.: 102:7145a,7148a

TITLE: Intramolecular Simmons-Smith reaction and other

synthetic alternatives to cyclopropanation of dienic

diazo ketones. Parallel decomposition pathways of a sterically congested diazo ketone and its vinylcyclopropane under thermal, photolytic,

acid-catalyzed, and radical-release

conditions

AUTHOR(S): Hudlicky, Tomas; Ranu, Brindaban C.; Naqvi, Saiyid M.;

Srnak, Ana

CORPORATE SOURCE: Dep. Chem., Virginia Polytechnic Inst. and State

Univ., Blacksburg, VA, 24061, USA

SOURCE: Journal of Organic Chemistry (1985), 50(1), 123-7

CODEN: JOCEAH; ISSN: 0022-3263

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 102:45503

GΙ

AB The decomposition of the diazo dienones I (R = H, Me; R1R2 = :N2) under thermal, photolytic, and acid catalyzed conditions was examined I (R = Me, R1R2 = :N2) was converted into geminal dihalides I (R1, R2 = Br, iodo), which were decomposed under conditions necessitating radical formation. The products, which were mostly the same, of these expts. included the cyclopropane II and the enone III. Reasons for the parallel outcomes of the expts. were discussed.